## \*Personal privacy information\*

Voluntary Industry Reporting Form for 6(a)(2) Adverse Effects Incident Information 2934

Provide all known, required information. If required data field information is unknown, designate as such in appropriate area Page 1 of 3

Row 1 Administrative	Reporter Name Submissible date.		Submission date.	Contact perso	on (if dif	ferent than reporter)	Internal ID 1939746-1
Data	Address  Sharps Chapel, TN 37866 USA			Address - OOH			
	Phone #	Phone #					
			date of incident el, TN	ate of incident Date registrant		Was incident part of larger study? No	
Row 2 Pesticide(s) Involved	EPA Registration # (Product 1) 62719-324		EPA Registration # (Product 2)		)	EPA Registration # (Product 3)	
	A.I. (s)		A.I. (s)	A.I. (s)			
	Product 1 name Rodeo Herbicide		Product 2 Name Method 24 SL Bayer			Product 3 Name Escort XP Bayer	
	Exposed to concentrate prior to dilution? No		Exposed to concentrate prior to dilution? No		)	Exposed to concentrate prior to dilution? No	
	Formulation Formulation			Formulation			
Row 3 Incident Circumstances	Evidence label directions were not followed? No Intentional misuse? No Applicator certified?	yard, school nursery/gree commercial woods, agric	enhouse, surface v turf, building/officultural (specify c ility, highway)).	vater, ice, forest/	Situation (act of using product): (example include mixing/loading, reentry, application transportation, repair/ maintenance of application equipment, manufacturing/ formulating).  See Incident Description Notes		
	UNK	Own Reside	nce				If no, why.
	How exposed: (examples include direct contact with treated surface, ingestion, spill, drift, runoff) See Incident Description Notes				.L.		11 11 8/1 /p.

Provide all known, required information. If required data field information is unknown, designate as such in appropriate area Page 2 of 3

#### Brief description of incident circumstances.

#### Nov 8 2016 9:40AM

Hx: Caller states property was sprayed on her property by a company spraying below an electrical line in late 06/16 The product was also sprayed on the well. The product was not intended to be sprayed on her garden. She was unaware the product was sprayed on the garden. She ate some of the berries within about 4 days of application. She became very ill. She developed nausea and diarrhea for 48 hours. She did not relate the illness to the product until a few days later when the bushes died. In early 07/2016 she developed insomnia. She saw her doctor for the insomnia on 09/20/2016. He prescribed Zolpidem for her insomnia. He did blood work and detected abnormal kidney function, She had an ultrasound on her kidneys, bladder and thyroid. The ultra sound shawed 4 small nodules on her thyroid. She has had fallow up blood work and her kidney function is still diminished. She has been tested Her primary wants her to see a nephrologist. She does not have an appointment scheduled at this time.

Her husband had a physical a few weeks after she did. He also had similar blood test results showing diminished kidney function.

Many other people in her area are having health problems, animals have died, bees died. She would like to know if the problems could be from mixing the products. The USDA has tested the wells and no contamination was found.

A: This product could cause some GI upset. But we would not be expected to cause any kidney problems. I am not able to address the other products used. We would not have information regarding the product being mixed with other companies products. Provided case number and told the caller if any new or unexpected symptoms develop or the symptoms are not improving ar resolving as we have discussed, please contact us 24/7 and refer to your reference number so that we can advise on further treatment or determine if referral to a healthcare professional might be needed.

Provide all known, required information. If required data field information is unknown, designate as such in appropriate area Page 3 of 3

Demographic information: Age: 59 Year(s) Sex: Female Occupation (if relevant) Not specified	Exposure route: Was adverse effect result suicide/homicide or attent suicide/homicide?			
If female, pregnant? NO	Was exposure occupational?  Not specified  If yes, days lost due to illness:  Not specified	Time between exposure and onset of symptoms:  I week or less		
Type of medical care sought: (examples include none, clinic, hospital emergency department, private physician, PCC, hospital inpatient). Non-emergent private MD/DVM	List signs/symptoms/adverse eff Gastrointestinal-Diarrhea Gastrointestinal-Emesis/Vomiti Genitourinary-diminished kidn	If lab tests were performed, list test names and results (If available, submit reports)  Creatinine .96 mg/dl  EGFR 58 and 59  mL/min/1.7		
Exposure data: NA Amount of pesticide: NA Exposure duration: Acute < 8hrs Patient weight: Unknown				
Human severity category: HC				

This box can be used to provide any explanatory or qualifying information surrounding the incident. (add additional pages if necessary)

Any relationship between the use of this product and the insidious development of the complications reported in this case is inconceivable and lacks biological plausibility. Secondly, the product use history is extremely vague and lacks any description of a known or defined point of direct exposure to this product. Even had cosual or incidental contact with this product occurred, such illness would be unexpected and is not consistent with the toxicological profile of this product.

Internal ID # 1939746-1

# \*Personal privacy information\*

Voluntary Industry Reporting Form for 6(a)(2) Adverse Effects Incident Information

Row I  Administrative Data	Reporter Name		Submission date.			ferent than reporter)	Internal ID 1939746-2
	Address  Sharps Chapel, TN 376 USA	Address Continuation of - O Phone #					
	Phone # (						
	Incident Status: Location and Sharps Chap USA 06/08/2016		1 date of incident net, TN	Date registrant became aware of incident.  11/08/2016		Was incident part of larger study? No	
Row 2  Pesticide(s) Involved	EPA Registration # (Product 1) 62719-324		EPA Registration # (Product 2)			EPA Registration # (Product 3)	
	A.I. (s)		A.I. (s)			A.I. (s)	
	Product 1 name Rodeo Herbicide		Product 2 Name Method 24 SL Bayer			Product 3 Name Escort XP Bayer	
	Exposed to concentrate prior to dilution? No		Exposed to concentrate prior to dilution? No		dilution? No		
	7 2 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7		Formulation	E 200 C 100 C 100 C 100 C		Formulation	
Row 3 Incident Circumstances	Evidence label directions were not followed? No Intentional misuse? No Applicator certified?	Incident site: (examples incluyard, school, industrial, nursery/greenhouse, surface commercial turf, building/off woods, agricultural (specify way (rail, utility, highway)).  Own Residence		include mixing/loading, reentry, apater, transportation, repair/ maintenance application equipment, manufacture		ntry, application, tenance of ufacturing/	
	UNK  How exposed: (examples include direct contact with treated surface. ingestion, spill, drift, runoff) See Incident Description Notes	Own Restat	me				

Provide all known, required information. If required data field information is unknown, designate as such in appropriate area. Page 2 of 3

#### Brief description of incident circumstances.

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Hx: Caller states property was sprayed on her property by a company spraying below an electrical line in late 06/16 The product was also sprayed on the well. The product was not intended to be sprayed on her garden. She was unaware the product was sprayed on the garden. She ate some of the berries within about 4 days of application. She became very ill. She developed nausea and diarrhea for 48 hours. She did not relate the illness to the product until a few days later when the bushes died. In early 07/2016 she developed insomnia. She saw her doctor for the insomnia on 09/20/2016. He prescribed Zolpidem for her insomnia. He did blood work and detected abnormal kidney function, She had an ultrasound on her kidneys, bladder and thyroid. The ultra sound showed 4 small nodules on her thyroid. She has had follow up blood work and her kidney function is still diminished. She has been tested Her primary wants her to see a nephrologist. She does not have an appointment scheduled at this time.

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Provide all known, required information. If required data field information is unknown, designate as such in appropriate area Page 3 of 3

Demographic information: Age: 60 Year(s) Sex: Male Occupation (if relevant) Not specified	Exposure route: Unknown route	Was adverse effect result of suicide/homicide or attempted suicide/homicide?	Was protective clothing worn (specify)? None Reported	
If female, pregnant?  NA	Was exposure occupational?  Not specified  If yes, days lost due to illness:  Not specified	Time between exposure and onset of symptoms:  Unable to determine		
Type of medical care sought: (examples include none, clinic, hospital emergency department, private physician. PCC, hospital inpatient). Non-emergent private MD/DVM	List signs/symptoms/adverse eff Heme/Hepatic-diminished kidn		If lab tests were performed, list test names and results (If available, submit reports)  None Reported	
Exposure data: NA Amount of pesticide: NA Exposure duration: Acute < 8hrs Patient weight: Unknown				
Human severity category: HC				

This box can be used to provide any explanatory or qualifying information surrounding the incident. (add additional pages if necessary)

Any relationship between the use of this product and the insidious development of the complications reported in this case is inconceivable and lacks biological plausibility. Secondly, the product use history is extremely vague and lacks any description of a known or defined point of direct exposure to this product. Even had casual or incidental contact with this product occurred, such illness would be unexpected and is not consistent with the toxicological profile of this product..

Internal ID # 1939/40-2